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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/705,227	11/12/2003	John D. Pruitt	029318-0985	3550
31049 7590 01/30/2009 Elan Drug Delivery, Inc. c/o Foley & Lardner 3000 K Street, N.W. Suite 500 Washington, DC 20007-5109			EXAMINER FUBARA, BLESSING M	
			ART UNIT 1618	PAPER NUMBER
			MAIL DATE 01/30/2009	DELIVERY MODE PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No. 10/705,227	Applicant(s) PRUITT ET AL.	
	Examiner BLESSING M. FUBARA	Art Unit 1618	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --
Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 23 October 2008.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-1 and 19-57 is/are pending in the application.
- 4a) Of the above claim(s) 27 and 31-36 is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-17, 19-26, 28-30 and 37-57 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. _____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- * See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|--|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date: _____ |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| Paper No(s)/Mail Date: _____ | 6) <input type="checkbox"/> Other: _____ |

DETAILED ACTION

Examiner acknowledges receipt of request for continued examination under 37 CFR

1.114, amendment and remarks filed 10/23/08. Claims 1, 19, 37 and 57 are amended. Claims 1-17 and 19-57 are pending. Claims 27 and 31-36 are withdrawn from consideration.

Response to Arguments

Previous rejections that are not reiterated herein are withdrawn.

Claim Rejections - 35 USC § 103

1. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

2. This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

3. Claims 1-17, 19-26 and 28-30 and 37-57 are rejected under 35 U.S.C. 103(a) as being unpatentable over Murakami et al. (US 6,287,596).

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Murakami discloses fast disintegrating compression molded product/tablet (abstract; column 4, lines 24-63; column 7, lines 11-15) comprising pharmaceutically active agents listed in column 5, line 61 to column 6, line 67 meeting the requirements of claims 1, 26, 30 and 56; for example, ibuprofen (column 6, line 3) is slightly or poorly water soluble meeting claim 52 and diphenhydramine hydrochloride (column 6, line 5) is soluble in water meeting claim 51, lubricants, diluents, coloring agents with the diluents being lactose or glucose or sucrose and the binders being acacia or pullulan or polyvinylpyrrolidone, flavoring agents, effervescent agents such as combinations of tartaric acid, malic acid and sodium carbonate or sodium bicarbonate (column 7, lines 13-45), and the pullulan and the effervescent couple meeting claims 1, 4, 5, 7-11, 14-16, 25 and 26 and 46. Claim 17 is the property of the product so that Murakami meets the claim. Friability is a property of the product; the disintegration time of the product of Murakami is in the order of seconds (column 9, lines 13-43) meeting claims 14 and 50; Murakami teaches that the rapidly disintegrating compression molded material is orally administered to infants and aged adults for the treatment of variety of diseases (column 9, lines 44-60; column 10, lines 34-40) meeting claim 37. Thus Murakami discloses a composition containing pullulan, effervescent couple, lactose and active agent. The composition of Murakami also contains surfactants (column 7, lines 43-49) meeting claims 21 and 28-30. Regarding claims 1, 19, 20 and 37 active agents are generally obtained in powder forms (column 5, lines 54 and 57) and the particles of the powder have sizes that meet the limitations of claims 18-20, with powder meeting the amorphous particle limitation of claim 55. The sugar alcohols in Murakami are namely D-mannitol, D-sorbitol, xylitol, maltitol, anhydrous maltose, hydrous maltose, anhydrous lactitol, hydrous lactitol, and reducing malt sugar syrup. Of these, D-

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mannitol, xylitol, and multitol (column 4, lines 54-58) and other excipients such as starches and celluloses (column 4, lines 29-53) meet claims 4-8, 11, 40, 41, 43, 44 and 47. Claim 48 is a product by process claim such that Murakami meets the claim. Regarding the amounts of the active agent and surface stabilizer, the claims 22-24 would have been obvious because the ordinary skilled artisan have the capabilities to use desired amounts of active agents and surface stabilizers in the composition Murakami for a rapidly disintegrating material. No specific particle size is recited in Murakami except that the active agents are in powder forms prior to inclusion in the dosage form such that the powder would have small particles sizes. Therefore, taking the general teaching of the reference, one having ordinary skill in the art at the time the invention was made would have reasonable expectation of success that using surface stabilizers and drugs in powder form and in appropriate and desired amounts would lead to solid dosage form that would have the anticipated rapid disintegration.

Response to Arguments

4. Applicant's arguments filed 10/23/08 have been fully considered but they are not persuasive.

Applicant argues that the claimed invention is distinguished over Murakami because the claimed invention does not contain erythritol while the Murakami's composition contains erythritol; that Murakami teaches that less than 30% by weight of erythritol leads to poor disintegration and dissolution, and that the claimed invention is a solid having friability of less than 1%. Applicant's arguments are not persuasive. a) erythritol is a sugar alcohol just as the sugar alcohols such sucrose, xylitol, mannose, sorbitol, trehalose, fructose contained in the invention (see at least instant claim 5). Therefore, the presence erythritol in the composition of

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Murakami would not materially affect the composition any more than the sugar alcohol recited in at least claim 5 would in the claimed composition. Thus, since the claimed composition/invention contains sugar alcohols, the consisting essentially of does not exclude any other sugar alcohols such as the erythritol. b) it is also noted that the sugar alcohols are present at about 1% to 99% such that even if Murakami desires to use greater than or equal to 30% erythritol, the $\geq 30\%$ falls within the claimed range (see at least claim 6) and the claims have not talked about using any amount of sugar alcohol. c) it is also noted that friability is a property of the dosage form and applicant has not factually shown that the dosage of Murakami is not friable. "When the PTO shows a sound basis for believing that the products of the applicant and the prior art are the same, the applicant has the burden of showing that they are not." In re Spada, 911 F.2d 705, 709, 15 USPQ2d 1655, 1658 (Fed. Cir. 1990).

5. No claim is allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to BLESSING M. FUBARA whose telephone number is (571)272-0594. The examiner can normally be reached on 7 a.m. to 5:30 p.m. (Monday to Thursday).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael G. Hartley can be reached on (571) 272-0616. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Blessing M. Fubara/
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